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Stephen Hordley

Dated

10 March 2003

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28 FEB 2002 E699605-1 002973
P01/7700 0.00-0204671.2

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AJC/P53051

2. Patent application number

(The Patent Office will fill in this part)

0204671.2

28 FEB 2002

3. Full name, address and postcode of the or of each applicant (underline all surnames)

British Nuclear Fuels Plc
Risley
WARRINGTON
Cheshire
WA3 6AS

Patents ADP number (if you know it)

350 108 001

If the applicant is a corporate body, give the country/state of its incorporation

UK

4. Title of the invention

ELECTROCHEMICAL CELL FOR METAL
PRODUCTION

5. Name of your agent (if you have one)

Harrison Goddard Foote

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

Belgrave Hall
Belgrave Street
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LS2 8DD

7631310002

Patents ADP number (if you know it)

14571001

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Country

Priority application number
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Date of filing
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7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(day / month / year)

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a) any applicant named in part 3 is not an inventor, or

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Description

7 ✓

Claim(s)

4 ✓

Abstract

1 ✓

Drawing(s)

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Request for preliminary examination and search (Patents Form 9/77)

1 ✓

Request for substantive examination (Patents Form 10/77)

Any other documents (please specify)

11.

I/We request the grant of a patent on the basis of this application.

Signature

Date

Harrison Goldard Doot

27 February 2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Tony Chalk

0113 233 0100

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ELECTROCHEMICAL CELL FOR METAL PRODUCTION

Field of the Invention

This invention relates to methods for the production of metals from oxides present in spent nuclear fuels and is particularly applicable to the production of actinides, specifically uranium. Methods of the present invention can be used in the treatment of irradiated fuels for producing actinides in metallic form suitable for use as feeds in subsequent electrorefining processes.

Background to the Invention

In the established art, two processes have been developed for the treatment of irradiated nuclear fuel making use of molten salts. As used herein, the term "molten salts" is intended to cover salts such as lithium chloride which melts at an elevated temperature and also ionic liquids which typically are liquid at room temperature or which melt at a temperature up to about 100°C.

The Dimitrovgrad SSC-RIAR process makes use of chemical oxidants (chlorine and oxygen gases) to react with powdered uranium dioxide fuel to form higher oxidation state compounds such as UO_2Cl_2 which are soluble in the molten salt. In an electrochemical cell the uranium compounds are reduced to UO_2 at the cathode, forming a dendritic deposit. This process has both technical and environmental limitations.

The second process, developed by the Argonne National Laboratory (ANL) is fundamentally an electrorefining technology which uses direct current to anodically oxidise uranium to form uranium ions in the molten salt electrolyte. At the cathode the uranium is reduced and electrodeposited as dendritic uranium metal.

The ANL process requires a metal feed. If oxide fuels are to be treated, it is necessary to reduce the uranium oxide (usually UO_2 pellets) to the metal. This reduction process is carried out chemically, using lithium metal in a LiCl or LiCl/KCl

molten salt, or a Li/Cd alloy, at 500 to 600°C. Alternatively, a salt transport process can be used involving a Cu-Mg-Ca alloy and molten CaCl_2 salt. However, in both reduction methods the by-products, Li_2O and CaO respectively, need to be recovered from the molten salt phase by an electrolysis step. Effectively this means a two stage process.

A disadvantage of the lithium reduction process for producing a metallic feed from an oxide is the production of Li_2O by-product. This requires recycle to make the process economic, and this is done by an electrolytic recovery of lithium metal.

10 Hence this is a two stage process, comprising a reduction step followed by a lithium recovery stage.

In co-pending PCT patent application WO 01/41152 there is disclosed a single step process for reducing to metallic form a metal oxide present in spent nuclear fuel, the process comprising cathodically electrolysing the oxide in the presence of a molten salt electrolyte, the potential of the cathode being controlled so as to favour oxygen ionisation over deposition of the metal from the cations present in the molten salt.

20 The process thereby involves the use of a single electrochemical process to reduce the metal oxide fuel to a metallic form, with oxygen, CO or CO_2 produced as the only by-products. The potential of the cathode is maintained and controlled so that only oxygen ionisation occurs and not the deposition of metal (eg Ca) from the cations (eg Ca ions) in the fused salt. Typically, the oxide comprises an actinide oxide, such as uranium oxide or irradiated uranium oxide.

25 The present inventors have, however, now effected an improvement to the process described in WO 01/41152 which allows a more practical, efficient and financially viable process to be adopted in the production of metals from oxides. The new process is particularly beneficial in that it facilitates the removal of bolted and

30 screwed fittings from the apparatus used for the production of the metal.

Statements of Invention

Thus, according to a first aspect of the present invention there is provided an apparatus for performing a process for reducing to metallic form metal oxides, the said apparatus being free from bolted or screwed fittings, and comprising an electrochemical cell which comprises a body or housing, a cathode container, and a cathode connector, wherein said body or housing is maintained as the cathode.

The body or housing of the cell which comprises the apparatus according to the first aspect of the invention is most conveniently maintained as the cathode by the provision of an electrical connection from a power supply, provided by means of a connector from the cathode terminal to the body or housing of the cell. Typically, said connector comprises a bolted connection, but this is positioned externally to the cell. No bolted or screwed connections are present within the cell, wherein a cathode connector is provided which is affixed to an internal surface of the cell, most preferably the base of the cell, generally by welding. Preferably said cathode connector comprises a cathode rail.

In operation, the body or housing of the cell is maintained as the cathode, and said cathode is brought into contact with the cathode container by means of the cathode connector. Thus, contact is made between the cathode container and the cathode connector in order to facilitate the electrolytic process. Contact may be most conveniently achieved by means of a simple press connection between the two components.

The cathode container preferably comprises a basket, such as a mesh basket, or vessel, typically a metal oxide retaining vessel, and – most preferably – comprises an assembly of such baskets or vessels. In order to effect electrical connection between such an assembly of cathode containers and the cell body during operation of the cell, it is necessary to provide a multiplicity of connectors and to effect contact between individual cathode baskets or vessels and individual connectors, preferably by means of a multiplicity of press connections.

Preferably the oxide is in contact with the cathode container and it is preferred that the cathode is in the form of a mesh basket or, most preferably, an assembly of mesh baskets, with the oxide being contained within the said baskets. In this case, contact between the assembly of cathode containers and the cathode connectors is most simply achieved when the cathode connectors are in the form of a multiplicity of cathode rails which are welded to the base of the cell, allowing press contact to be brought about by the weight of the oxide feedstock in the cathode basket. This represents the most preferred embodiment of the first aspect of the invention. The anode may be any suitable inert anode, such as carbon.

10 According to a second aspect of the present invention, there is provided a process for reducing to metallic form metal oxides, the process comprising cathodically electrolysing the oxide in the presence of a molten salt electrolyte in an apparatus according to the first aspect of the invention, the potential of the cathode being controlled so as to favour oxygen ionisation over deposition of metal from the cations present in the molten salt.

The molten salt electrolyte may be any suitable molten salt or mixture of such salts, for instance chloride salts, preferably CaCl_2 and/or BaCl_2 .

20 Preferably, the oxide treated by the process according to the second aspect of the invention comprises an oxide present in spent nuclear fuel. Typically, the oxide comprises an actinide oxide, such as uranium oxide or irradiated uranium oxide, or mixed uranium/plutonium oxides. The uranium oxide is commonly uranium dioxide.

25 Alternatively, the oxide may comprise the oxide of a metal such as zirconium or hafnium.

In such a process, wherein the oxide comprises an oxide present in spent nuclear fuel, the fuel may be first treated mechanically to remove its zircaloy cladding before it is added to the electrolytic cell. Alternatively, the zircaloy cladding may be treated with

30

the fuel. The fuel may require to be sheared into sections of small length prior to treatment in order to expose the oxide fuel to the molten salt.

5 Said oxide may be in any physical form, and this is generally dependent on the particular chemical nature of the spent nuclear fuel and the processing to which the material has previously been subjected. For example, the fuel may comprise a powder, an amorphous mass, or a dense solid agglomerate. In any event, the material may be treated according to the method of the second aspect of the present invention by connection to an electrical circuit such that it serves as the cathode during
10 electrolysis; as previously disclosed, connection to the circuit may be conveniently effected by the use of a cathode basket, into which the material is placed.

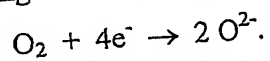
The process provides a single electrochemical process to reduce the metal oxide fuel to a metallic form, with oxygen produced as the only by-product. The potential of
15 the cathode is maintained and controlled so that only oxygen ionisation occurs and not the deposition of metal (eg Ca) from the cations (eg Ca ions) in the fused salt.

The process according to the second aspect of the invention provides a more practical, efficient and financially viable means for the production of metal from
20 oxides. The electrical continuity of the system which is provided is effective since all the components are held in the cathodic potential region and are, therefore, metallic in nature.

Detailed Description of the Invention

25 In order to carry out an embodiment of the process according to the second aspect of the present invention, using a metal oxide which comprises spent nuclear fuel, an electrolytic cell is assembled which has a carbon anode, a mesh basket cathode and a cathode rail connector welded to the base of the cell. Irradiated oxide fuel is placed in the mesh basket. The electrolyte consists of a molten salt or a mixture of such
30 salts comprising, for example, chloride salts such as CaCl_2 or BaCl_2 . A voltage is applied between the cathode and the anode. At the cathode the reaction involves the

diffusion of oxygen atoms to the surface of the solid, followed by ionisation according to the reaction:



5 The oxide ions which are produced dissolve in the electrolyte and are transferred to the anode where they are re-oxidised to produce oxygen, CO or CO₂ gases. The potential at the cathode may be controlled, via a third reference electrode, to ensure that the reaction occurring at the cathode is oxygen ionisation and not deposition of eg Ca metal from the cations in the fused salt. Electrolysis at elevated temperatures
10 ~~results in an increased rate of oxygen diffusion, thereby also encouraging ionisation~~ rather than metal deposition.

After electrolysis the irradiated fuel is left in the form of a metallic solid at the cathode. This metallic solid, which contains fission products, can be removed or used directly as the feed for an electrorefining process. The remaining components
15 of the cell may be re-used immediately without the need for any cleaning.

In an alternative embodiment in accordance with the present invention the electrolytic ionisation of oxygen and the electrorefining processes are carried out in the same cell and the same salt system. In such cases a second cathode, or cathode
20 assembly, may be required.

It is to be emphasised that the advantage of the apparatus and process of the present invention is that they facilitate the performance of an effectively single stage process which provides advantages over the prior art in terms of practicality, efficiency and
25 cost. The process is advantageously used for the treatment of irradiated oxide nuclear fuel, possibly in the form of pellets and, most particularly, is applied to fuels such as uranium oxide, and mixed uranium and plutonium fuels.

In order to fully illustrate the above and other features of the apparatus according to
30 the first aspect of the invention, without in any way limiting the scope of the

invention, a preferred embodiments will now be described with reference to the drawing.

Description of the Drawing

5 Figure 1 shows an apparatus according to the first aspect of the invention which comprises a series of electrolytic cells, the bodies and housings of which are, in use, connected by means of cathode rails 1 to cathode baskets 2. The cells also comprise anodes 4, surrounded by anode protective sleeves 3, and connected to an electrical supply by means of electrical connections 6, these being protected by insulation
10 shrouds 5.

In operation, the cathode baskets 2, which are equipped with basket handles 8, cooled by means of cooling chambers 7, are loaded with spent nuclear fuel pellets and lowered into the electrolyte 9 to a point at which contact is made with cathode rails 1
15 and, thereby, with the cell housing, which is maintained as the cathode, thereby allowing processing to commence.

CLAIMS

1. An apparatus for performing a process for reducing to metallic form metal oxides, the said apparatus being free from bolted or screwed fittings and comprising an electrochemical cell which comprises a body or housing, a cathode container, and a cathode connector, wherein said body or housing is maintained as the cathode.
2. An apparatus as claimed in claim 1 wherein said cathode connector is affixed to an internal surface of the cell.
3. An apparatus as claimed in claim 2 wherein said cathode connector is affixed to an internal surface of the cell by means of welding.
4. An apparatus as claimed in any one of claims 1, 2 or 3 wherein an electrical connection from the cathode container to the body or housing of the cell is provided by means of a cathode connector.
5. An apparatus as claimed in claim 4 wherein said electrical connection is provided by means of a press connection.
6. An apparatus as claimed in any preceding claim wherein said cathode connector comprises a cathode rail.
7. An apparatus as claimed in claim 6 wherein said rail is welded to the base of the cell.
8. An apparatus as claimed in any preceding claim wherein the cathode container comprises a mesh basket or metal oxide retaining vessel.

9. An apparatus as claimed in any preceding claim wherein the cathode container comprises an assembly of cathode containers.
10. An apparatus as claimed in claim 9 wherein said assembly comprises an assembly of mesh baskets or metal oxide retaining vessels.
11. An apparatus as claimed in any preceding claim wherein the anode is a carbon anode.
12. An apparatus as claimed in any preceding claim wherein the body or housing of the cell is maintained as the cathode by the provision of an electrical connection from a power supply to the body or housing of the cell.
13. A process for reducing to metallic form metal oxides, the process comprising cathodically electrolysing the oxide in the presence of a molten salt electrolyte in an apparatus as claimed in any one of claims 1 to 12, the potential of the cathode being controlled so as to favour oxygen ionisation over deposition of metal from the cations present in the molten salt.
14. A process as claimed in claim 13 wherein the body or housing of the cell is maintained as the cathode and is brought into contact with the cathode container by means of a press connection between said container and a cathode connector.
15. A process as claimed in claim 13 or 14 wherein the oxide comprises the oxide of zirconium or hafnium.
16. A process as claimed in claim 13 or 14 wherein the oxide comprises an oxide present in spent nuclear fuel.

17. A process as claimed in claim 16 wherein the oxide comprises an actinide oxide.

18. A process as claimed in claim 17 wherein the actinide oxide comprises uranium oxide, irradiated uranium oxide or mixed uranium/plutonium oxide fuel pellets.

19. A process as claimed in claim 18 wherein the uranium oxide comprises uranium dioxide.

20. A process as claimed in any of claims 13 to 19 wherein the oxide is located in a mesh basket which forms the cathode.

21. A process as claimed in any of claims 13 to 20 wherein the molten salt electrolyte comprises at least one chloride salt.

22. A process as claimed in claim 21 wherein the chloride salt is CaCl_2 or BaCl_2 .

23. A process as claimed in any one of claims 16 to 22 wherein the fuel is treated together with its cladding.

24. A process as claimed in any one of claims 16 to 22 wherein the cladding is removed from the fuel prior to treatment.

25. A process as claimed in any of claims 13 to 24 wherein the metal resulting from the process is used as the feed for an electrorefining process.

26. A process as claimed in claim 25 wherein the electrorefining process is carried out in the same electrolytic cell as the electrolytic reduction process.

27. An apparatus according to claim 1 and substantially as herein described.

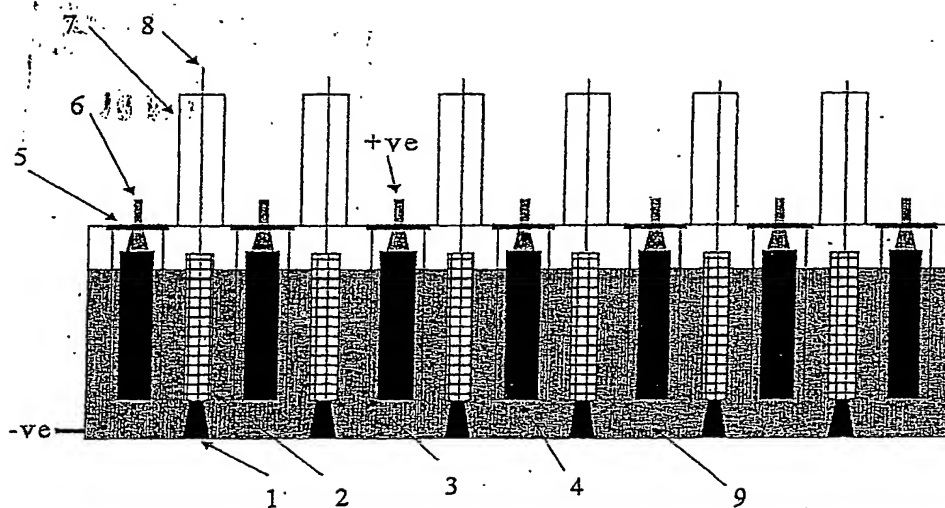
28. A process according to claim 13 and substantially as herein described.

ABSTRACT

The invention provides a process for reducing to metallic form metal oxides, the process comprising cathodically electrolysing the oxide in the presence of a molten salt electrolyte whilst controlling the potential of the cathode so as to favour oxygen ionisation over deposition of metal from the cations present in the molten salt, and an apparatus for performing the said process, the said apparatus being free from bolted or screwed fittings and comprising an electrochemical cell which comprises a body or housing, a cathode container, and a cathode connector, wherein said body or

housing is maintained as the cathode. The process provides a single electrochemical process to reduce the metal oxide fuel to a metallic form, with oxygen produced as the only by-product, and offers a more practical, efficient and financially viable means for the production of metal from oxides than is available from the prior art. The metallic solid which is produced can be removed or used directly as the feed for an electrorefining process. In an alternative embodiment, the electrolytic ionisation of oxygen and the electrorefining processes are carried out in the same cell and the same salt system.

P53051.2



- 1 Weld cathode rail
- 2 Cathode basket
- 3 Anode protective sleeve
- 4 Anode
- 5 Insulation shroud
- 6 Anode electrical connection
- 7 Cooling chamber
- 8 Basket handle
- 9 Electrolyte

FIGURE 1

1/1

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